WHAT IS CLAIMED IS:

1 1. A compound having a formula that is a member selected from:

$$R^{6} \xrightarrow{Q} Q \xrightarrow{R^{1}} Q \xrightarrow{R^{2}} Q \xrightarrow{R^{2}} Q \xrightarrow{R^{2}} Q \xrightarrow{R^{4}} Q \xrightarrow{R^{3}} Q \xrightarrow{R^{4}} Q \xrightarrow{R^{4}$$

3 wherein

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- 4 R^1 is H, CH_2OR^7 , $COOR^7$ or OR^7
- 5 in which
- R⁷ represents H, substituted or unsubstituted alkyl or substituted or unsubstituted heteroalkyl;
- R² is a member selected from H, OH, an activating group and a moiety that includes a nucleotide;
- 10 R³, R⁴, R⁵, R⁶ and R⁶ are independently selected from H, substituted or unsubstituted alkyl, OR⁹, and NHC(O)R¹⁰
- 12 wherein
- 13 R⁹ and R¹⁰ are independently selected from H, substituted or unsubstituted alkyl or substituted or unsubstituted heteroalkyl,
- and at least one of R³, R⁴, R⁵, R⁶ and R⁶ includes a polymeric modifying moiety.
- 1 2. The compound according to claim 1 wherein \mathbb{R}^2 has the formula:

- 3 in which R⁸ is a nucleoside.
- 1 3. The compound according to claim 2 wherein R⁸ is a member selected from cytosine,
- 2 uridine, guanosine, adenosine and thymidine.

1 4. The compound according to claim 1 wherein at least one of R³, R⁴, R⁵ and R⁶ includes
2 the moiety:

$$(R^{11})_{w} - L - \xi.$$

- 4 wherein
- 5 R¹¹ is a polymeric modifying moiety;
- 6 L is a member selected from a bond and a linking group; and
- 7 w is selected from the integers from 1 to 6.
- 1 5. The compound according to claim 4 wherein said linking group is a member selected
- 2 from substituted or unsubstituted alkyl and substituted or unsubstituted heteroalkyl moieties.
- 1 6. The compound according to claim 5 wherein the moiety:

3 has the formula:

$$R^{12}-X^2$$
 C
 X^4
 R^{13}
 X^4

5 wherein

- X^2 and X^4 are independently selected from linkage fragments;
- 7 X^a is a linkage fragment;
- 8 R¹² and R¹³ are independently selected polymeric arms; and
- 9 c is an integer from 1 to 20.
- 1 7. The compound according to claim 5 wherein said linking group has the formula:

$$\xi \xrightarrow{\qquad} X^a \xrightarrow{\qquad} L^1 \xrightarrow{\qquad} X^b \xrightarrow{\qquad} \xi$$

- 3 in which
- 4 X^a and X^b are independently selected linkage fragments; and
- 5 L¹ is a member selected from a bond, substituted or unsubstituted alkyl or substituted or unsubstituted heteroalkyl.

- 1 8. The compound according to claim 7 wherein X^a and X^b are linkage fragments
- 2 independently selected from S, SC(O)NH, HNC(O)S, SC(O)O, O, NH, NHC(O), (O)CNH
- and NHC(O)O, and OC(O)NH.
- 1 9. The compound according to claim 5 wherein said linker comprises an acyl moiety.
- 1 10. The compound according to claim 9 wherein L-R¹¹ has the formula:

$$\xi$$
—NHC(O)(CH₂)_s—NHC(O)—R¹¹

- 3 in which
- 4 s is an integer from 0 to 20; and
- 5 R¹¹ is said polymeric modifying moiety.
- 1 11. The compound according to claim 1, wherein said polymeric modifying moiety has
- 2 the formula:

$$R^{12}-X^{2}$$
 $X^{5}-C-\xi$
 $R^{13}-X^{4}$

4 wherein

- 5 X^2 and X^4 are independently selected from linkage fragments;
- 6 X⁵ is a non-reactive group; and
- R¹² and R¹³ are independently selected polymeric arms.
- 1 12. The compound according to claim 11 wherein X^2 and X^4 are linkage fragments
- 2 independently selected from S, SC(O)NH, HNC(O)S, SC(O)O, O, NH, NHC(O), (O)CNH
- 3 and NHC(O)O, OC(O)NH and (CH₂) $_{g}Y$ "
- 4 wherein
- 5 g is an integer from 1 to 50; and
- 6 Y" is a member selected from O, S and NH.
- 1 13. The compound according to claim 11 wherein
- 2 X⁴ is a peptide bond; and
- 3 R¹³ is an amino acid residue.
- 1 14. The compound according to claim 1 having the formula:

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3 in which

D is a member selected from -OH and $(R^{11})_{w'}$ -L-;

G represents is a member selected from H, $(R^{11})_{w'}$ -L- and -C(O)(C₁-C₆)alkyl;

6 w' is an integer from 2 to 6, and

at least one of D and G is $(R^{11})_{w'}$ -L-.

1 15. The compound according to claim 14 having the formula:

$$R^{12}-X^{2}$$
 $X^{5}-C$
 L^{3}
 $R^{13}-X^{3}$
 R^{5}
 R^{4}
 $R^{12}-X^{2}$
 $R^{12}-X^{2}$
 $R^{12}-X^{2}$
 $R^{12}-X^{2}$
 $R^{12}-X^{2}$
 $R^{12}-X^{2}$
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 $R^{13}-X^{3}$

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wherein

L^a is a member selected from substituted or unsubstituted alkyl and substituted or unsubstituted heteroalkyl.

1 16. The compound according to claim 1 having the formula:

$$R^{12}-X^{2}$$
 $X^{5}-C$
 L^{a}
 $R^{13}-X^{4}$
 R^{5}
 R^{1}
 R^{2}
 R^{3}
 R^{3}
 R^{4}
 R^{4}
 R^{3}
 R^{4}
 R^{4}
 R^{5}
 R^{4}
 R^{5}
 R^{4}
 R^{5}
 R^{5}
 R^{6}
 R^{7}
 R^{7}

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wherein

L^a is a member selected from an amino acid residue and a peptidyl residue having from 2 to 4 amino acid residues;

6 X² and X⁴ are independently selected from linkage fragments;

- 7 X⁵ is a non-reactive group; and
- 8 R¹² and R¹³ are independently selected polymeric arms
- 1 17. The compound according to claim 16 having the formula:

3 wherein

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- 4 X^2 and X^4 are independently selected from linkage fragments;
- 5 X^a is a linkage fragment;
- 6 R¹² and R¹³ are independently selected polymeric arms; and
- 7 c is an integer from 1 to 20.
- 1 18. The compound according to claim 1, having the formula:

3 wherein

- 4 AA-NH is an amino acid residue; and
- P is a polymeric modifying group.
- 1 19. The compound according to claim 18 wherein -AA-NH is -CH₂NH.
- 1 20. The compound according to claim 1 wherein said compound is a substrate for an
- 2 enzyme that transfers a sugar moiety from a member selected from an activated sugar, a
- 3 nucleotide sugar and combinations thereof onto an acceptor moiety of a substrate.
- 1 21. The compound according to claim 20 wherein said acceptor moiety is a member
- 2 selected from a glycosyl residue, an amino acid residue and an aglycone.

- 1 22. A method of preparing cytidine monophosphate sialic acid-poly(ethylene glycol), said 2 method comprising:
 - (a) contacting mannosamine with an activated, N-protected amino acid under conditions appropriate to form an amide conjugate between said mannosamine and the N-protected amino acid;

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- (b) contacting said amide conjugate with pyruvate and sialic acid aldolase under conditions appropriate to convert said amide conjugate to a sialic acid amide conjugate;
- (c) contacting said sialic acid amide conjugate with cytidine triphosphates, and a synthetase under conditions appropriate to form a cytidine monophosphate sialic acid amide conjugate;
 - (d) removing the N-protecting group from said cytidine monophosphate sialic acid amide conjugate, thereby producing a free amine; and
- 13 (e) contacting said free amine with an activated PEG, thereby forming said 14 cytidine monophosphate sialic acid-poly(ethylene glycol).
- 1 23. The method according to claim 21, wherein said activated N-protected amino acid has 2 the formula: